Oral plasma zinc tolerance test in patients with protein energy malnutrition

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SUMMARY Zinc absorption was measured in 37 children with malnutrition using the oral zinc tolerance test (22.5 mg elementary zinc) and the results compared with those of a group of healthy control subjects. The increase in plasma zinc was significantly lower in patients with marasmic kwashiorkor than in the control group. The zinc tolerance test was, however, normal in marasmic patients.

We conclude that zinc deficiency occurs in some types of protein energy malnutrition, and that malabsorption may aggravate zinc deficiency. It is reasonable to give higher doses of zinc than are usually recommended during oral zinc supplementation in patients with protein energy malnutrition.

Protein energy malnutrition is still an important problem in many parts of the world. Certain clinical features are common to patients with protein energy malnutrition and those with zinc deficiency, including diarrhoea, anorexia, growth retardation, muscular atrophy, and a tendency to infections. This similarity has led many workers to study zinc absorption in patients with protein calorie malnutrition. Some found that plasma zinc concentrations were low, and others found decreased serum zinc concentrations in malnutrition, together with skin lesions.

Protein energy malnutrition is associated with various degrees of intestinal changes, which include gastrointestinal mucosal atrophy and structural alterations in the jejunal villi. Apart from inadequate dietary intake, digestive and absorptive defects may be important in the aetiology of protein energy malnutrition. It is, however, difficult to say whether these defects are causes or effects of the malnutrition.

The aim of this study was to determine whether there is a decrease in intestinal absorption of zinc in protein energy malnutrition in addition to pre-existing zinc deficiency and malabsorption.

Subjects and methods

Thirty seven children (19 boys and 18 girls) aged between 5 months and 5 years with protein energy malnutrition were compared with 10 healthy children (eight boys and two girls) aged between 6 months and 5 years. The patients were classified according to the McLaren system.

Routine laboratory tests were performed using standard methods. Serum alkaline phosphatase activities were measured (Sigma Diagnostics), and plasma and erythrocyte zinc concentrations were assessed by an atomic absorption spectrophotometer (Perkin Elmer Model 403). Blood samples (4.5 ml) were collected using zinc free materials and anticoagulated with zinc free oxalate solution. Plasma was separated by centrifugation and the plasma zinc concentration determined using 3:1 dilution of plasma. Erythrocyte zinc concentrations were determined by the method of Rosney and Gorfren.

For the oral plasma zinc tolerance test blood was taken from subjects to determine fasting plasma zinc after an eight hour overnight fast. A dose of 100 mg hydrated zinc sulphate (22.5 mg elementary zinc) was then given. The subjects were allowed only water during the test. After the oral test dose of zinc, blood was drawn at one, two and four hours for measurement of plasma zinc concentrations. Using the data obtained, the increases in plasma zinc concentrations above fasting values were plotted. The control group was tested in the same way.

Mean (SD) values were calculated for all measurements. Student’s t test was used to assess the significance of differences between the groups.
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**Results**

Thirty seven malnourished children aged 5 months to 5 years and 10 healthy control subjects aged 6 months to 5 years were studied. Thirteen children had marasmus, 18 marasmic kwashiorkor, and six kwashiorkor.

Diarrhoea was present in 22 (60%), oedema in five (14%), anal, gluteal, and oral dermatitis in 19 (52%), and hair loss in 26 (70%).

Height (p<0.05), weight (p<0.01), mid arm circumference (p<0.001), haemoglobin concentration (p<0.01), total protein and albumin (p<0.01), and globulin (p<0.05) concentrations, and alkaline phosphatase activity (p<0.01) were significantly decreased in the malnourished group compared with the control group (tables 1 and 2). The decreases in serum albumin concentrations were especially prominent in the patients with marasmic kwashiorkor and kwashiorkor (p<0.05 and p<0.01, respectively).

Plasma zinc and erythrocyte zinc concentrations and the results of the oral plasma zinc tolerance tests are shown in table 3. Plasma zinc and erythrocyte zinc concentrations were significantly lower in every subgroup compared with the control group (p<0.01).

When patients with marasmic kwashiorkor and kwashiorkor were examined together after the zinc test dose had been given, the increase in plasma zinc was significantly less in the second hour compared with the control group (p<0.05). On the other hand, the increase in plasma zinc was normal in the subgroup of marasmic patients (figure).

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**Table 1 Clinical and laboratory findings in 37 patients with protein energy malnutrition and 10 control subjects. Values are expressed as mean (SD)**

<table>
<thead>
<tr>
<th></th>
<th>Patients</th>
<th>Control subjects</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Height (cm)</td>
<td>68.8 (13.4)</td>
<td>78.0 (9.9)</td>
<td>0.001</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>6.7 (3.2)</td>
<td>10.0 (3.2)</td>
<td>0.01</td>
</tr>
<tr>
<td>Mid arm circumference (cm)</td>
<td>10.4 (2.2)</td>
<td>15.4 (1.2)</td>
<td>0.05</td>
</tr>
<tr>
<td>Haemoglobin concentration (g/l)</td>
<td>97 (20)</td>
<td>117 (10)</td>
<td>0.01</td>
</tr>
<tr>
<td>Alkaline phosphatase activity (IU/l)</td>
<td>60.7 (38.8)</td>
<td>101.7 (39.0)</td>
<td>0.01</td>
</tr>
<tr>
<td>Total protein concentration (g/l)</td>
<td>55 (10)</td>
<td>69 (5)</td>
<td>NS</td>
</tr>
<tr>
<td>Albumin concentration (g/l)</td>
<td>35 (9)</td>
<td>43 (5)</td>
<td>0.01</td>
</tr>
<tr>
<td>Globulin concentration (g/l)</td>
<td>20 (4)</td>
<td>25 (7)</td>
<td>0.01</td>
</tr>
</tbody>
</table>

*Compared with normal subjects.

**Table 2 Protein concentrations (g/l) in 37 patients with protein energy malnutrition. Values are expressed as mean (SD)**

<table>
<thead>
<tr>
<th></th>
<th>Marasmus (n=13)</th>
<th>Marasmic kwashiorkor (n=18)</th>
<th>p* Value</th>
<th>Kwashiorkor (n=6)</th>
<th>p* Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total protein</td>
<td>61 (9)</td>
<td>54 (8)</td>
<td>0.01</td>
<td>43 (10)</td>
<td>0.001</td>
</tr>
<tr>
<td>Albumin</td>
<td>39 (8)</td>
<td>35 (8)</td>
<td>0.05</td>
<td>26 (8)</td>
<td>0.001</td>
</tr>
<tr>
<td>Globulin</td>
<td>22 (4)</td>
<td>20 (4)</td>
<td>0.001</td>
<td>17 (4)</td>
<td>0.005</td>
</tr>
</tbody>
</table>

**Table 3 Zinc concentrations in 37 patients with protein energy malnutrition and 10 control subjects**

<table>
<thead>
<tr>
<th></th>
<th>Patients</th>
<th>Marasmus (n=13)</th>
<th>Marasmic kwashiorkor (n=18)</th>
<th>Kwashiorkor (n=6)</th>
<th>Total (n=37)</th>
<th>Control subjects (n=10)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean (SD) fasting plasma zinc (μmol/l)</td>
<td>13.9 (4-6)*</td>
<td>15.6 (3-7)</td>
<td>13.0 (2-9)</td>
<td>14.6 (3-8)**</td>
<td>18.4 (3-9)</td>
<td>19.9 (12-1-24.5)</td>
</tr>
<tr>
<td>Median (range)</td>
<td>13.8 (6-9-19-9)</td>
<td>15.1 (6-9-22-2)</td>
<td>12.3 (10-2-16-7)</td>
<td>14.4 (6-9-22-2)</td>
<td>17.1 (6-0)</td>
<td>18.5 (10-4-23-9)</td>
</tr>
<tr>
<td>Mean (SD) erythrocyte zinc (mg/l)</td>
<td>11.9 (4-3)*</td>
<td>8.0 (2-7)**</td>
<td>8.8 (0-8)**</td>
<td>9.6 (3-7)**</td>
<td>18.5 (10-4-23-9)</td>
<td></td>
</tr>
<tr>
<td>Median (range)</td>
<td>11.4 (5-4-18-9)</td>
<td>8.2 (4-1-12-9)</td>
<td>9.4 (8-2-18-1)</td>
<td>9.1 (4-1-21-7)</td>
<td>18.5 (10-4-23-9)</td>
<td></td>
</tr>
<tr>
<td>Mean (SD) increase in plasma zinc (μmol/l):</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>At one hour</td>
<td>11.4 (5-8)</td>
<td>4.8 (4-6)**</td>
<td>5.8 (5-3)</td>
<td>7.7 (5-7)*</td>
<td>12.1 (7-6)</td>
<td>17.5 (6-5)</td>
</tr>
<tr>
<td>At two hours</td>
<td>18.2 (11-0)</td>
<td>10.4 (6-8)*</td>
<td>8.0 (7-6)*</td>
<td>12.1 (9-2)</td>
<td>17.5 (6-5)</td>
<td>2.2 (4-5)</td>
</tr>
<tr>
<td>At four hours</td>
<td>6.3 (5-6)</td>
<td>3.4 (4-2)</td>
<td>2.5 (3-6)</td>
<td>4.3 (5-6)</td>
<td>2.2 (4-5)</td>
<td></td>
</tr>
</tbody>
</table>

*p<0.05; **p<0.01, and ***p<0.001 compared with control subjects.
The mitosis rate in the cells of the gastrointestinal, immune and haematopoietic systems is high. The effects of zinc deficiency as a result of the important action of zinc on protein synthesis, therefore, are observed mostly in these systems, so it is reasonable to claim that the chronic zinc deficiency in protein energy malnutrition might lead to pathological changes in the intestinal mucosa.

Many alterations in the gastrointestinal tracts of malnourished children have been reported. A genetic zinc deficiency disease, acrodermatitis enteropathica, causes severe intestinal mucosal atrophy, and zinc supplementation is effective in returning the mucosal to normal.

To measure intestinal zinc absorption the oral plasma zinc tolerance test was conducted in patient and control groups. The oral plasma zinc tolerance test, which measures the increase in plasma zinc concentrations, has been widely used as an assessment of intestinal zinc absorption. Under standard conditions the test gives valuable information about the absorption of zinc by the small intestine. Valberg et al compared radioactive zinc absorption with the oral plasma zinc tolerance test and found that the tests gave comparable results. In both our subgroups of patients (marasmic kwashioror and kwashiorkor) the plasma zinc curve (figure) was significantly depressed compared with that of the control group.

In patients with marasmus, the plasma zinc tolerance test was normal, confirming the results of Brunser et al, who studied jejunal biopsy specimens of patients with marasmus and found normal mucosa.

We conclude that zinc deficiency in patients with protein energy malnutrition can induce intestinal mucosal changes that may aggravate the zinc deficiency, thus creating a vicious circle. It is therefore reasonable to give higher doses of zinc than are usually recommended when giving oral zinc supplementation to patients with protein energy malnutrition.

References

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